



**[1,2,4]TRIAZOLO[1,5-c]PYRIMIDINE DERIVATIVES****Publication number:** WO0017201**Publication date:** 2000-03-30**Inventor:** SHIMADA JUNICHI (JP); IMMA HIRONORI (JP);  
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SHIOZAKI SHIZUO (JP); KANDA TOMOYUKI (JP);  
KUWANA YOSHIHISA (JP)**Classification:****- International:** **A61P25/24; A61P25/28; C07D487/04; A61P25/00;**  
**C07D487/00; (IPC1-7): C07D487/04; A61K31/505****- European:** C07D487/04**Application number:** WO1999JP05176 19990922**Priority number(s):** JP19980267178 19980922**Also published as:** EP1116722 (A1)  
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**Cited documents:** WO9842711  
 US5270311**Report a data error here****Abstract of WO0017201**

[1,2,4]Triazolo[1,5-c]pyrimidine derivatives represented by general formula (I) or pharmacologically acceptable salts thereof which show adenosine A2A receptor antagonism, wherein R<1> represents heteroaryl, etc.; R<2> represents hydrogen, etc.; na and nb represent each an integer of 0 to 4; Q represents hydrogen, etc.; R<6> represents hydrogen, etc.; R<3> represents (i) hydroxy, (ii) hydroxy (lower alkyl), (iii) lower alkoxy, or (iv) imidazo[1,2-a]pyridyl, etc.; and R<4> and R<5> represent each lower alkyl or aryl, or R<4> and R<5> form together with the adjacent carbon atom a saturated carbon ring when R<3> is any of (i) to (iii); or R<4> and R<5> represent each hydrogen, lower alkyl or aryl, or R<4> and R<5> form together with the adjacent carbon atom a saturated carbon ring when R<3> is (iv).

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